Serial No.: 10/044,848 Group Art Unit No.: 1615 Preliminary Amendment B

Amendments to the claims

1-19 (Cancelled)

- (Currently Amended) A process to prepare pharmaceutical tablets containing paroxetine, on a commercial scale, which process comprises the steps of:
 - a) dry admixing paroxetine and dry excipients in a mixer to form a mixture; or
- b) dry admixing paroxetine and dry excipients, compressing the resulting combination into a slug material or roller compacting the resulting combination into a strand material, and milling the prepared material into a free flowing mixture; and
- c) compressing the mixture into tablets; provided that the excipients include at least one of; sodium starch glycollate, dicalcium phosphate and magnesium stearate.
- 21. (Currently Amended) A process to prepare pharmaceutical tablets containing paroxetine, on a commercial scale, which process comprises the steps of:
 - dry admixing paroxetine and dry excipients in a mixer to form a mixture; or
- dry admixing paroxetine and dry excipients, compressing the resulting combination into a slug material or roller compacting the resulting combination into a strand material, and milling the prepared material into a free flowing mixture; and
- c) compressing the mixture into tablets;
 provided that the excipients include at least one of; sodium starch glycollate, dicalcium phosphate and magnesium stearate;
 and further provided that one of the excipients that is compressed into tablets is not microcrystalline cellulose.
- 22. (Previously Presented) A process according to claim 20 in which the amount of paroxetine in each tablet is selected from: 10 mg, 20 mg, 30 mg, 40 mg and 50 mg, wherein the amount of paroxetine is expressed as the free base.
- 23. (Previously Presented) A process according to claim 21 in which the amount of paroxetine in each tablet is selected from: 10 mg, 20 mg, 30 mg, 40 mg and 50 mg, wherein the amount of paroxetine is expressed as the free base.
- 24. (Previously Presented) A process according to claim 22 in which the amount of paroxetine in each tablet is about 10 mg, wherein the amount of paroxetine is expressed as the free base.



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- 25. (Previously Presented) A process according to claim 23 in which the amount of paroxetine in each tablet is about 10 mg, wherein the amount of paroxetine is expressed as the free base.
- 26. (Previously Presented) A process according to claim 22 in which the amount of paroxetine in each tablet is about 20 mg, wherein the amount of paroxetine is expressed as the free base.
- 27. (Previously Presented) A process according to claim 23 in which the amount of paroxetine in each tablet is about 20 mg, wherein the amount of paroxetine is expressed as the free base.
- 28. (Previously Presented) A process according to claim 22 in which the amount of paroxetine in each tablet is about 30 mg, wherein the amount of paroxetine is expressed as the free base.
- 29. (Previously Presented) A process according to claim 23 in which the amount of paroxetine in each tablet is about 30 mg, wherein the amount of paroxetine is expressed as the free base.
- 30. (Previously Presented) A process according to claim 22 in which the amount of paroxetine in each tablet is about 40 mg, wherein the amount of paroxetine is expressed as the free base.
- (Previously Presented) A process according to claim 23 in which the amount of paroxetine in each tablet is about 40 mg, wherein the amount of paroxetine is expressed as the free base.
- 32. (Previously Presented) A process according to claim 22 in which the amount of paroxetine in each tablet is about 50 mg, wherein the amount of paroxetine is expressed as the free base.
- 33. (Previously Presented) A process according to claim 23 in which the amount of paroxetine in each tablet is about 50 mg, wherein the amount of paroxetine is expressed as the free base.

